phenylene group optionally substituted by a second fluorine, chlorine or bromine atom or by a second $C_{1-3}$-alkyl group,

a thienylene, thiazolylene, pyridinylene, pyrimidinylene, pyrazinylene or pyridazinylene group optionally substituted in the carbon skeleton by a $C_{1-3}$-alkyl group,

$R_1$ denotes a $C_{1-3}$-alkyl group optionally substituted by an amino, $C_{1-3}$-alkylamino, di-($C_{1-3}$-alkyl)-amino, phenyl, naphthyl, heteroaryl or 4- to 7-membered cycloalkyleneimino group,

a $C_{3-7}$-cycloalkyl group which is substituted in the 1 position by a 5- to 7-membered cycloalkyleneiminocarbonyl group,

an amino, $C_{1-5}$-alkylamino, $C_{5-7}$-cycloalkylamino or phenyl-$C_{1-3}$-alkylamino group which may in each case be substituted at the amino-nitrogen atom by a benzoyl or phenylsulphonyl group or by a $C_{1-3}$-alkyl or $C_{1-3}$-alkylcarbonyl group optionally substituted in the $C_{1-3}$-alkyl moiety by a carboxy group,

a 4- to 7-membered cycloalkyleneiminocarbonyl or cycloalkyleneiminosulphonyl group optionally substituted by a $C_{1-3}$-alkyl group,

an aminosulphonyl group optionally substituted by one or two $C_{1-3}$-alkyl groups, a phenyl group optionally substituted by a fluorine, chlorine or bromine atom, by a trifluoromethyl, aminosulphonyl, $C_{1-3}$-alkyl or $C_{1-3}$-alkoxy group, which may additionally be substituted by a fluorine, chlorine or bromine atom or by a trifluoromethyl, $C_{1-3}$-alkyl or $C_{1-3}$-alkoxy group,

a $C_{1-3}$-alkoxy, phenyl-$C_{1-3}$-alkoxy, heteroaryloxy or heteroaryloxy-$C_{1-3}$-alkoxy group wherein the alkoxy moiety may be substituted in the 2 or 3 position in each case by an amino, $C_{1-3}$-alkylamino or di-($C_{1-3}$-alkyl)-amino group,

a $C_{3-7}$-cycloalkoxy group, wherein the methylene group in the 3 or 4 position in a $C_{5-7}$-cycloalkoxy group may be replaced by an -NH group, and said -NH group may be optionally substituted

by a $C_{1-3}$-alkyl group which may be substituted in the 2 or 3 position by an amino, $C_{1-3}$-alkylamino or di-($C_{1-3}$-alkyl)-amino group, by a $C_{1-3}$-alkylcarbonyl, arylcarbonyl or arylsulphonyl group or

by an aminocarbonyl, $C_{1-3}$-alkylaminocarbonyl or di-($C_{1-3}$-alkyl)-aminocarbonyl group, wherein in each case the oxygen atom of the carbonyl group is replaced by an imino group,

$R_2$ denotes a hydrogen, fluorine, chlorine or bromine atom, a $C_{1-3}$-alkyl, hydroxy or $C_{1-3}$-alkoxy group,

$R_3$ denotes a hydrogen atom or a $C_{1-3}$-alkyl group,

$R_4$ denotes a hydrogen atom or a $C_{1-3}$-alkyl group optionally substituted by a carboxy group and

$R_5$ denotes a cyano group or an amidino group optionally substituted by one or two $C_{1-3}$-alkyl groups, said heteroaryl groups consisting of a 5-membered heteroaryl group optionally substituted by a $C_{1-3}$-alkyl group which contains in the heteroaromatic moiety,

an imino group optionally substituted by a $C_{1-3}$-alkyl group, or an oxygen or sulphur atom,

an imino group optionally substituted by a $C_{1-3}$-alkyl group and an oxygen, sulphur or nitrogen atom,

an imino group optionally substituted by a $C_{1-3}$-alkyl group and two nitrogen atoms or

an oxygen or sulphur atom and two nitrogen atoms,

or a 6-membered heteroarylene group optionally substituted by a $C_{1-3}$-alkyl group which contains one or two nitrogen atoms in the heteroaromatic moiety,

an isomer or salt thereof.

2. (amended) The compound of formula I according to claim 1 wherein

one of the groups m or n denotes the number 0 and
the other group m or n denotes the number 1,

Ar denotes a phenylene group optionally substituted by a fluorine, chlorine or bromine atom or by a methyl, hydroxy, methoxy or benzyloxy group, which may be substituted by another methyl group,

$R_1$ denotes a phenyl group optionally substituted by a fluorine, chlorine or bromine atom or by a trifluoromethyl, aminosulphonyl, $C_{1-3}$-alkyl or $C_{1-3}$-alkoxy group, which may additionally be substituted by a fluorine, chlorine or bromine atom or by a trifluoromethyl, $C_{1-3}$-alkyl or $C_{1-3}$-alkoxy group,

a methyl group substituted by a dimethylamino, pyrrolidino or imidazolyl group, wherein the imidazolyl moiety may be substituted by a methyl group,

an amino, $C_{1-5}$-alkylamino, cyclopentylamino or benzylamino group which may be substituted at the amino-nitrogen atom by a carboxy-$C_{1-2}$-alkyl, $C_{1-3}$-alkoxycarbonyl-$C_{1-2}$-alkyl, carboxy-$C_{1-2}$-alkylcarbonyl or $C_{1-3}$-alkoxycarbonyl-$C_{1-2}$-alkylcarbonyl group,

a benzoylamino or phenylsulphonylamino group,

a cyclopropyl group which is substituted in the 1 position by a 5- to 7-membered cycloalkyleneiminocarbonyl group,

an optionally methyl-substituted pyrrolidinocarbonyl, piperidinocarbonyl, pyrrolidinosulphonyl or piperidinosulphonyl group,

a $C_{1-3}$-alkoxy group wherein the alkoxy moiety in the 2 or 3 position may be substituted in each case by an amino, $C_{1-3}$-alkylamino or di-($C_{1-3}$-alkyl)-amino group,

a phenyl-$C_{1-3}$-alkoxy or pyridinyloxy group,

a $C_{5-7}$-cycloalkoxy group wherein the methylene group in the 3 or 4 position may be replaced by an -NH group, said -NH group may be substituted

by a $C_{1-3}$-alkyl or $C_{2-3}$-alkanoyl group,

A2

by a $C_{2-3}$-alkanoyl or aminocarbonyl group wherein in each case the oxygen atom of the carbonyl group is replaced by an imino group,

$R_2$ denotes a hydrogen, fluorine, chlorine or bromine atom, a methyl, hydroxy or methoxy group,

$R_3$ denotes a hydrogen atom or a methyl group,

$R_4$ denotes a hydrogen atom or a methyl or ethyl group optionally substituted by a carboxy or $C_{1-3}$-alkoxycarbonyl group and

$R_5$ denotes a cyano group or an amidino group optionally substituted by a $C_{1-6}$-alkoxycarbonyl or benzoyl group,

or an isomer or salt thereof.

3. (amended) The compounds of formula I according to claim 1, wherein

one of the groups m or n denotes the number 0 and
the other group m or n denotes the number 1,

Ar denotes a phenylene group optionally substituted by a methyl, hydroxy, methoxy or benzyloxy group,

$R_1$ denotes a phenyl group optionally substituted by a fluorine, chlorine or bromine atom or by a trifluoromethyl, aminosulphonyl, $C_{1-3}$-alkyl or $C_{1-3}$-alkoxy group, which may additionally be substituted by a fluorine, chlorine or bromine atom or by a trifluoromethyl, $C_{1-3}$-alkyl or $C_{1-3}$-alkoxy group,

a cyclopropyl group which is substituted in the 1 position by a 5- to 7-membered cycloalkyleneiminocarbonyl group, or a 4- to 7-membered cycloalkyleneiminocarbonyl group,

an optionally methyl-substituted pyrrolidinocarbonyl, piperidinocarbonyl or pyrrolidinosulphonyl group,

$R_2$ denotes a hydrogen, fluorine, chlorine or bromine atom or a methyl group,

$R_3$ denotes a hydrogen atom or a methyl group,

$R_4$ denotes a hydrogen atom or a methyl or ethyl group substituted by a carboxy, methoxycarbonyl or ethoxycarbonyl group and

$R_5$ denotes an amidino group optionally substituted by a $C_{1-6}$-alkoxycarbonyl or benzoyl group,

or an the isomers or the salts thereof.

4. (amended) A compound of the formula I according to claim 1 selected from the following compounds:

(a) 2-(5-carbamimidoyl-2-hydroxy-phenyl)-N-[3-methyl-4-(pyrrolidin-1-yl-carbonyl)-phenyl]-acetamide,

(b) 2-(2-benzyloxy-5-carbamimidoyl-phenyl)-N-(2-ethoxycarbonyl-ethyl)-N-[3-methyl-4-(pyrrolidin-1-yl-carbonyl)-phenyl]-acetamide,

(c) 2-(2-hydroxy-5-carbamimidoyl-phenyl)-N-(2-ethoxycarbonylethyl)-N-[3-methyl-4-(pyrrolidin-1-yl-carbonyl)-phenyl]-acetamide,

(d) 2-(2-hydroxy-5-carbamimidoyl-phenyl)-N-(2-carboxy-ethyl)-N-[3-methyl-4-(pyrrolidin-1-yl-carbonyl)-phenyl]-acetamide,

(e) 2-(5-carbamimidoyl-2-hydroxy-phenyl)-N-[3-methyl-4-(piperidin-1-yl-carbonyl)-phenyl]-acetamide and

(f) 2-(5-carbamimidoyl-2-hydroxy-phenyl)-N-[3-methyl-4-(2-aminosulphonyl-phenyl)-phenyl]-acetamide,

wherein the amidino group may additionally be substituted by a $C_{1-6}$-alkoxycarbonyl or benzoyl group, and the salts thereof.

5. (amended) A compound of formula 1 according to claim 1 as follows: 2-(5-Carbamimidoyl-2-hydroxy-phenyl)-N-[3-methyl-4-(pyrrolidin-1-yl-carbonyl)-phenyl]-acetamide and the salts thereof.

6. (amended) A pharmaceutical composition comprising a compound according to claim 1 or a physiologically acceptable salt thereof according to claim 1 wherein $R_5$ denotes said amidino groups.

7. (amended) Pharmaceutical compositions containing a compound according to claim 1, wherein $R_5$ denotes said amidino groups.

8. (amended) A method of treating a patient in need of a pharmaceutical composition having an antithrombotic activity or factor Xa inhibiting activity by administering to said patient a therapeutically effective amount of a component according to claim 1 wherein $R_5$ denotes said amidino groups.